

The strongest inverse correlation was observed between ADC-mean and either SUV-mean or -SUV-max in tumor lesions.

References:

(1) Functional imaging of head and neck squamous cell carcinoma with diffusion-weighted MRI and FDG PET/CT: quantitative analysis of ADC and SUV. Varoquaux A, Rager O, Lovblad KO, Masterson K, Dulguerov P, Ratib O, Becker CD, Becker M.

587

Increasing tumor grade separability by combining MR parameters

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Purpose/Introduction: MR diffusion, perfusion and spectroscopic data provide complementary information in brain tumor grading. We show that combining MR parameters of different modalities can improve diagnostic accuracy.

Subjects and Methods: MR data: MR data were acquired on a 3T Philips Achieva scanner from 5 glioma patients: 2 grade II astrocytomas, 1 grade II oligodendroglioma, 2 grade IV glioblastomas. DSC-MRI was acquired with a GE-EPI sequence: TR/TE=1350/30msec, with 60 dynamic scans during the first pass of a 0.1mmol/kg body weight bolus of Dotarem (Guerbet) injected at 4ml/sec. Cerebral Blood Volume (CBV) values were calculated using DPTTools (www.fmrtools.org). DKI data were acquired using an SE-EPI sequence: TR/TE=3200/90msec, $\delta/J=20/48.3$ msec, $b=700,1000,2800$ sec/mm². Mean Diffusion (MD) and Mean Kurtosis (MK) values were calculated as in [1]. MRSI data were acquired using PRESS volume selection, TR/TE=2000/35msec, MOIST water suppression, receiver bandwidth=2000Hz. AQSES-MRSI [2] was used for metabolite quantification. All imaging modalities were coregistered to anatomical data.

Analysis: Perfusion and diffusion maps were normalized with respect to NAWM. 5 parameters were considered: CBV, MD, MK, Lips/TCho (Lipids over total Choline) and Cre/sum (Creatine over summed metabolites). All voxels in the active tumor ROIs were included, resulting in class C1 of 1821 low-grade glioma voxels and class C2 of 873 high-grade glioma voxels (Fig.1).

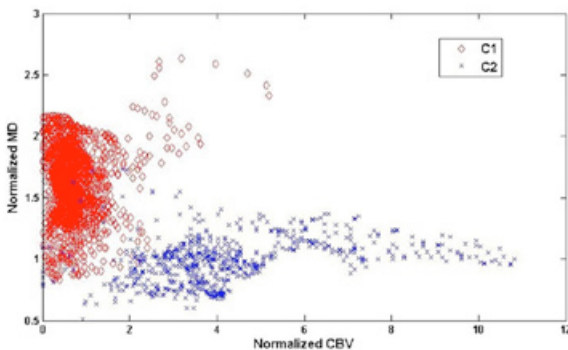


Fig.1: Scatter plot of CBV versus MD for LGG class C1 and HGG class C2

Intra-class Homogeneity and inter-class Separation were computed:

$$Hom_{C_i} = \frac{1}{N_{C_i}} \sum_k D(x_k, \mu_{C_i})$$

$$Sep_{C_i, C_j} = D(\mu_{C_i}, \mu_{C_j})$$

with N_{C_i} the number of data points in class C_i , D a distance measure, x_k the k^{th} point in class C_i and μ_{C_i} the class centroid. Mahalanobis is used as distance measure, taking into account the directional spread of the data. Based on Hom and Sep , the HS-index is calculated:

$$HS_{C_i, C_j} = Sep_{C_i, C_j} - Hom_{C_i}$$

Results: Table1 and Table2 show HS-indices of C1 with respect to C2 and vice-versa. The HS-values on the diagonal are for 1 parameter only. Green cells indicate the combinations for which the HS-index is increased with respect to both parameters' individual HS-values. The HS-value when combining all parameters is shown at the bottom of the table.

	Lips/TCho	Cre/sum	CBV	MD	MK
Lips/TCho	5.98	6.29	9.31	5.67	5.64
Cre/sum		0.59	6.30	1.09	0.71
CBV			6.64	6.80	6.45
MD				1.48	1.15
MK					0.44
All params	15.51				

Table 1: HS_{C1,C2} values for combined MR parameters

	Lips/TCho	Cre/sum	CBV	MD	MK
Lips/TCho	1.01	2.04	1.22	2.66	1.40
Cre/sum		2.09	2.73	3.03	2.44
CBV			1.01	3.66	1.87
MD				2.72	2.69
MK					1.33
All params	6.67				

Table 2: HS_{C2,C1} values for combined MR parameters

Discussion/Conclusion: The separability of tumor grade classes can improve by combining certain MR parameters. However, a slight decrease of the combined HS-index is found for other combinations. For all parameters combined, the HS-index is higher than for any combination of 2 parameters. Tumor grade separability can be maximized by considering the optimal combination of parameters.

References:

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- Croitor Sava A.R. et al, NMR Biomed. 2011 Aug;24(7):824-35.

588

Longitudinal T1 mapping for the follow-up of patients with abdominal solid tumors

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Purpose/Introduction: Dynamic MRI with contrast injection is generally used to characterize the micro vasculature in patients with abdominal solid tumors. Response Evaluation Criteria in Solid Tumors (RECIST) [1] are conventional morphological criteria to evaluate the tumor response to a treatment. This work proposes a new approach based on the T1 mapping to quantify the tumor response.

Subjects and Methods: Seven patients with abdominal solid tumors were examined using a dedicated MRI protocol before and during the course of